California Drug Recall Information

Recall Name
Hospira Recalls One Lot Of Dobutamine Injection, USP, 250 MG, 20 ML, Single-Dose Fliptop Vial Due to Visible Particulate

Recall Date	Product Description	Recalling Firm	Recall Reason
01/10/14	Dobutamine Injection, USP, 250 MG, 20 ML, Single-Dose Fliptop Vial **NDC # 0409-2344-02**	**Hospira, Inc.** Lake Forest, IL	*Due to a confirmed customer report of discolored solution.* *Upon review of the complaint, a chip in the glass neck of the vial was identified, as well as glass particulates within the solution.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
N/A	Suspect Lot: • **Lot 27-352-DK** Product Photo	**CA**, nationwide	Distributed from August 2013 through September 2013.

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm397339.htm